

FORM PTO-1390 (Modified) (REV 11-2000)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NUMBER	
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371				213301US0PCT	
				U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 09/926168)	
INTERNATIONAL APPLICATION NO. PCT/EP00/02416		INTERNATIONAL FILING DATE 17 March 2000		PRIORITY DATE CLAIMED 19 March 1999	
TITLE OF INVENTION PROCESS FOR THE PREPARATION OF SURFACTANT ALCOHOLS AND SURFACTANT ALCOHOL ETHERS, THE PREPARED PRODUCTS AND THEIR USE					
APPLICANT(S) FOR DO/EO/US MAAS Heiko et al.					
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:					
<ol style="list-style-type: none"> 1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. 2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. 3. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (24) indicated below. 4. <input checked="" type="checkbox"/> The US has been elected by the expiration of 19 months from the priority date (Article 31). 5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371 (c) (2)) <ol style="list-style-type: none"> a. <input type="checkbox"/> is attached hereto (required only if not communicated by the International Bureau). b. <input checked="" type="checkbox"/> has been communicated by the International Bureau. c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). 6. <input checked="" type="checkbox"/> An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)). <ol style="list-style-type: none"> a. <input checked="" type="checkbox"/> is attached hereto. b. <input type="checkbox"/> has been previously submitted under 35 U.S.C. 154(d)(4). 7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3)) <ol style="list-style-type: none"> a. <input type="checkbox"/> are attached hereto (required only if not communicated by the International Bureau). b. <input type="checkbox"/> have been communicated by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input checked="" type="checkbox"/> have not been made and will not be made. 8. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). 9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)). 10. <input type="checkbox"/> An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)). 11. <input type="checkbox"/> A copy of the International Preliminary Examination Report (PCT/IPEA/409). 12. <input checked="" type="checkbox"/> A copy of the International Search Report (PCT/ISA/210). <p>Items 13 to 20 below concern document(s) or information included:</p> <ol style="list-style-type: none"> 13. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. 14. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. 15. <input checked="" type="checkbox"/> A FIRST preliminary amendment. 16. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment. 17. <input type="checkbox"/> A substitute specification. 18. <input type="checkbox"/> A change of power of attorney and/or address letter. 19. <input type="checkbox"/> A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825. 20. <input type="checkbox"/> A second copy of the published international application under 35 U.S.C. 154(d)(4). 21. <input type="checkbox"/> A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4). 22. <input type="checkbox"/> Certificate of Mailing by Express Mail 23. <input checked="" type="checkbox"/> Other items or information: <p>Request for Consideration of Documents Cited in International Search Report Notice of Priority PCT/IB/304</p>					

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR **09/926168**) INTERNATIONAL APPLICATION NO. **PCT/EP00/02416** ATTORNEY'S DOCKET NUMBER **213301US0PCT**

24. The following fees are submitted:


BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :

<input type="checkbox"/> Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO	\$1000.00	CALCULATIONS PTO USE ONLY		
<input checked="" type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO	\$860.00			
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO	\$710.00			
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4)	\$690.00			
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4)	\$100.00			
ENTER APPROPRIATE BASIC FEE AMOUNT =		\$860.00		
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).		\$0.00		
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	
Total claims	21 - 20 =	1	x \$18.00	\$18.00
Independent claims	1 - 3 =	0	x \$80.00	\$0.00
Multiple Dependent Claims (check if applicable). <input type="checkbox"/>				\$0.00
TOTAL OF ABOVE CALCULATIONS =				\$878.00
<input type="checkbox"/> Applicant claims small entity status. (See 37 CFR 1.27). The fees indicated above are reduced by 1/2.				\$0.00
SUBTOTAL =				\$878.00
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).				\$0.00
TOTAL NATIONAL FEE =				\$878.00
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable). <input type="checkbox"/>				\$0.00
TOTAL FEES ENCLOSED =				\$878.00
				Amount to be refunded \$
				charged \$

- a. A check in the amount of **\$878.00** to cover the above fees is enclosed.
- b. Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees. A duplicate copy of this sheet is enclosed.
- c. The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. **15-0030**. A duplicate copy of this sheet is enclosed.
- d. Fees are to be charged to a credit card. **WARNING:** Information on this form may become public. **Credit card information should not be included on this form.** Provide credit card information and authorization on PTO-2038.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:



22850

Surinder Sachar
Registration No. 34,428

Surinder Sachar

SIGNATURE _____

Norman F. Oblon

NAME _____

24,618

REGISTRATION NUMBER _____

Sept. 18 2001

DATE _____

09/926168

531 Rec'd PCT/ 18 SEP 2001

213301US-0PCT

IN THE UNITED STATES PATENT & TRADEMARK OFFICE

IN RE APPLICATION OF :
HEIKO MAAS ET AL : ATTN: APPLICATION DIVISION
SERIAL NO: NEW US PCT APPLN :
(Based on PCT/EP00/02416) :
FILED: HEREWITH :
FOR: PROCESS FOR THE PREPARATION:
OF SURFACTANT ALCOHOL ETHERS,
THE PREPARED PRODUCTS AND
THEIR USE

PRELIMINARY AMENDMENT

ASSISTANT COMMISSIONER FOR PATENTS
WASHINGTON, D.C. 20231

SIR:

Prior to examination on the merits, please amend the above-identified application as follows:

IN THE CLAIMS

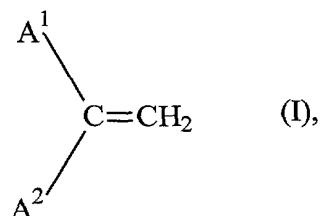
Please cancel Claims 14-18.

Please amend the claims as shown on the marked-up copy following this amendment to read as follows:

3. (Amended) A process as claimed in claim 1, wherein process step a), the dimerization, is carried out with heterogeneous catalysis.
4. (Amended) A process as claimed in claim 1, wherein a dimerization catalyst is used which comprises at least one element of subgroup VIII of the Periodic Table and the

09/926168

catalyst composition and the reaction conditions are chosen such that a dimer mixture is obtained which comprises less than 10% by weight of compounds which have a structural element of the formula I (vinylidene group)



in which A¹ and A² are aliphatic hydrocarbon radicals.

6. (Amended) An olefin mixture as claimed in claim 5, which comprises a proportion of branched components greater than 85%, and an unbranched olefin proportion below 15%.

7. (Amended) An olefin mixture as claimed in claim 5, wherein predominantly groups having (y-4) and (y-5) carbon atoms are bonded to the branching sites of the main chain, where y is the number of carbon atoms in the dimerized monomer.

8. (Amended) An olefin mixture as claimed in claim 5, wherein the branched components of the dimerization mixture, in the region of 1/4 to 3/4 of the chain length of the main chain, have a branch, or two branches on adjacent carbon atoms.

9. (Amended) An olefin mixture as claimed in claim 5, wherein predominantly groups having one or two carbon atoms are bonded to the branching sites of the main chain.

10. (Amended) An olefin mixture as claimed in claim 5, wherein, in the case of the branched components, the ratio of aliphatic to olefinic hydrogen atoms is in the range

$$H_{\text{aliph.}} : H_{\text{olefin.}} = 47 : 1 \text{ to } 11 : 1.$$

11. (Amended) An olefin mixture as claimed in claim 5, wherein, in the case of the branched components, the ratio of aliphatic to olefinic hydrogen atoms is in the range

$$H_{\text{aliph.}} : H_{\text{olefin.}} = 23 : 1 \text{ to } 14 : 1.$$

Please add new Claims 19-26.

19. (New) An olefin mixture as claimed in claim 8, wherein the branched components of the dimerization mixture, in the region of 1/3 to 2/3 of the chain length of the main chain, have a branch, or two branches on adjacent carbon atoms.

20. (New) The olefin mixture claimed in claim 6 wherein the branched olefins comprise a proportion greater than 90%.

21. (New) The olefin mixture claimed in claim 6 wherein the unbranched olefins comprise a proportion less than 10%.

22. (New) A nonionic surfactant comprising the surfactant alcohol alkoxylation product of claim 12.

23. (New) A method for the preparation of a surfactant comprising chemically modifying the surfactant alcohol or alkoxylation product thereof of claim 12.

24. (New) A method for the preparation of alkanol glycoside and polyglycoside mixtures comprising single or multiple reaction (glycosidation, polyglycosidation) of the surfactant alcohols of claim 12 with mono-, di- or polysaccharides with the exclusion of water and with acid catalysis or with O-acetylsaccharide halides.

25. (New) A method for the preparation of surface-active sulfates comprising esterification of the surfactant alcohols and alkoxylation products thereof of claim 12 with sulfuric acid or sulfuric acid derivatives to give acidic alkyl sulfates or alkyl ether sulfates.

26. (New) A method for the preparation of surface-active phosphates comprising esterification of the surfactant alcohols and alkoxylation products thereof of claim 12 with phosphoric acid or its derivatives to give acidic alkyl phosphates or alkyl ether phosphates.

REMARKS

Claims 1-13 and 19-26 are active in the present application. Claims 1-13 have been amended to remove multiple dependencies. Claims 14-18 have been cancelled and replaced by new Claims 22-26 that conform to U.S. Patent Practice. New Claims 19-21 are supported by original Claims 6 and 8. No new matter is added. An action on the merits and allowance of claims is solicited.

Respectfully submitted,

OBLON, SPIVAK, McCLELLAND,
MAIER & NEUSTADT, P.C.



Norman F. Oblon
Attorney of Record
Registration No. 24,618

Daniel J. Pereira, Ph.D.
Registration No. 45,518

Tel.: (703) 413-3000
Fax: (703) 413-2220
NFO:DJP:SUKOS\la
I:\atty\SUKOS\213301US-PR.WPD

213301US-0PCT

<p>Marked-Up Copy</p> <p>Serial No: _____</p> <p>Amendment Filed on: _____</p> <p style="text-align: center;">9-18-01</p>

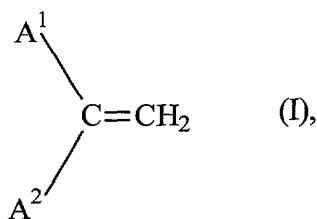
IN THE CLAIMS

Claims 14-18 (Cancelled).

Please amend the claims as follows:

--3. (Amended) A process as claimed in [at least one of claims 1 and 2] claim 1, wherein process step a), the dimerization, is carried out with heterogeneous catalysis.

4. (Amended) A process as claimed in [at least one of claims 1 to 3] claim 1, wherein a dimerization catalyst is used which comprises at least one element of subgroup VIII of the Periodic Table and the catalyst composition and the reaction conditions are chosen such that a dimer mixture is obtained which comprises less than 10% by weight of compounds which have a structural element of the formula I (vinylidene group)



in which A¹ and A² are aliphatic hydrocarbon radicals.

6. (Amended) An olefin mixture as claimed in claim 5, which comprises a proportion of branched components [-] greater than 85%[, in particular greater than 90% - of

components with branches], and an unbranched olefin [a low proportion - below 15, in particular below 10% - of unbranched olefins] proportion below 15%.

7. (Amended) An olefin mixture as claimed in [at least one of claims 5 and 6] claim 5, wherein predominantly groups having (y-4) and (y-5) carbon atoms are bonded to the branching sites of the main chain, where y is the number of carbon atoms in the dimerized monomer.

8. (Amended) An olefin mixture as claimed in [at least one of claims 5 to 7] claim 5, wherein the branched components of the dimerization mixture, in the region of 1/4 to 3/4[, preferably from 1/3 to 2/3,] of the chain length of the main chain, have a branch, or two branches on adjacent carbon atoms.

9. (Amended) An olefin mixture as claimed in [at least one of claims 5 to 8] claim 5, wherein predominantly groups having one or two carbon atoms are bonded to the branching sites of the main chain.

10. (Amended) An olefin mixture as claimed in [at least one of claims 5 to 9] claim 5, wherein, in the case of the branched components, the ratio of aliphatic to olefinic hydrogen atoms is in the range

$$H_{\text{aliph.}} : H_{\text{olefin.}} = 47 : 1 \text{ to } 11 : 1.$$

11. (Amended) An olefin mixture as claimed in [at least one of claims 5 to 10] claim 5, wherein, in the case of the branched components, the ratio of aliphatic to olefinic hydrogen atoms is in the range

$$H_{\text{aliph.}} : H_{\text{olefin.}} = 23 : 1 \text{ to } 14 : 1.--$$

Claims 19-26 (New).

Linear olefins of suitable chain length are currently accessible mainly by two processes:

In the Fischer-Tropsch synthesis, as well as paraffins, olefin isomer mixtures form as coupling products.

- 5 The oligomerization of ethylene has established itself as a further source for obtaining suitable olefins on an industrial scale. In this process, the catalysts used are alkylaluminums and also homogeneous nickel catalysts, as in the case of the known SHOP process from Shell (Weissermel/Arpe, Industrielle organische Chemie [Industrial Organic Chemistry]).
- 10 Olefin fractions of suitable chain length are further processed to give surfactant alcohols. The use of ethylene has the disadvantage of high feed material costs for the monomer building block. Processes for the preparation of surfactants which are based on ethylene as starting material are therefore important economically.
- 15 For the dimerization of olefins, a number of processes are known. For example, the reaction can be carried out over a heterogeneous cobalt oxide/carbon catalyst (FR-A-1 403 273), in the presence of acids such as sulfuric or phosphoric acid (FR 964 922), with an alkylaluminum catalyst (WO 97/16398), or with a homogeneously dissolved nickel complex catalyst (US-A-4 069 273). According to
- 20 the details in US-A-4 069 273, the use of these nickel complex catalysts (the complexing agent used being 1,5-cyclooctadiene or 1,1,1,5,5,5-hexafluoropentane-2,4-dione) gives highly linear olefins with a high proportion of dimerization products.
- FR-A-1 274 529 describes the Lewis-acid-catalyzed dimerization of
- 25 methylpentenes, where the Lewis acid used is boron trifluoride. This process has the disadvantage that it is difficult to separate off the catalyst from the reaction product. As a result, not only are products contaminated with catalyst residues obtained, but the catalyst loss is also considerable.
- 30 Functionalization of the olefins to give alcohols with extension of the carbon skeleton about a carbon atom advantageously takes place via the hydroformylation reaction, which gives a mixture of aldehydes and alcohols, which can then be hydrogenated to give alcohols. Approximately 7 million metric tons of products per annum are produced worldwide using the hydroformylation of olefins. An
- 35 overview of catalysts and reaction conditions for the hydroformylation process is given, for example, by Beller et al. in Journal of Molecular Catalysis, A104 (1995),

17-85 and also in Ullmann's Encyclopedia of Industrial Chemistry, vol. A5 (1986), page 217 et seq., page 333, and the relevant literature references.

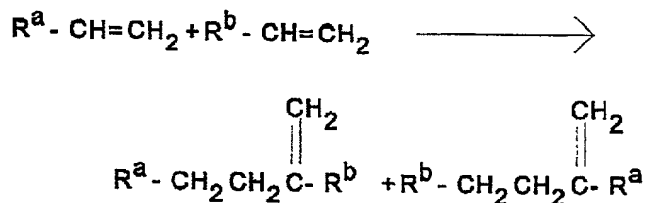
5 From WO 98/23566 it is known that sulfates, alkoxyates, alkoxyulfates and
carboxylates of a mixture of branched alkanols (oxo alcohols) exhibit good surface
activity in cold water and have good biodegradability. The alkanols in the mixture
used have a chain length of greater than 8 carbon atoms, having on average from
0.7 to 3 branches. The alkanol mixture can be prepared, for example by
10 hydroformylation, from mixtures of branched olefins which for their part can be
obtained either by skeletal isomerization or by dimerization of internal, linear
olefins.

A given advantage of the process is that no C₃- or C₄-olefin stream is used for the
preparation of the dimerization feed. It follows from this that, according to the
current prior art, the olefins subjected to dimerization therein must have been
15 prepared from ethylene (e.g. SHOP process). Since ethylene is a relatively
expensive starting material for surfactant manufacture, ethylene-based processes
have a disadvantage in terms of cost compared with processes which start from C₃-
and/or C₄-olefin streams.

20 The structure of the components of the oxo alkanol mixture depends on the type of
olefin mixture which has been subjected to hydroformylation. Olefin mixtures
which have been obtained by skeletal isomerization from alpha-olefin mixtures
lead to alkanols which are branched predominantly at the ends of the main chain,
i.e. in positions 2 and 3, calculated from the end of the chain in each case.

25 The surface-active end products are obtained from the alkanol mixtures either by
oxidation of the -CH₂OH group to give the carboxyl group, or by sulfation of the
alkanols or their alkoxyates.

Similar processes for the preparation of surfactants are described in the PCT Patent
30 Application WO 97/38957 and in EP-A-787 704. Also in the processes described
therein, an alpha-olefin is dimerized to give a mixture of predominantly vinylidene-
branched olefin dimers:



5 The vinylidene compounds are then double-bond-isomerized, such that the double bond migrates from the end of the chain further into the center, and are then subjected to hydroformylation to give an oxo alcohol mixture. The latter is then further reacted, e.g. by sulfation to give surfactants. A serious disadvantage of this process is that it starts from alpha-olefins. Alpha-olefins are obtained, for example, by transition-metal-catalyzed oligomerization of ethylene, Ziegler build-up reaction, wax cracking or Fischer-Tropsch processes and are therefore relatively expensive starting materials for the manufacture of surfactants. A further
10 considerable disadvantage of this known surfactant preparation process is that a skeletal isomerization must be inserted in the process between the dimerization of the alpha-olefins and the hydroformylation of the dimerization product if predominantly branched products are desired. Because it uses a starting material which is relatively expensive for surfactant manufacture and because of the need to insert an additional process step, the isomerization, this known process is at a
15 considerable disadvantage in terms of cost.

US-A-5,780,694 describes the preparation and use of alcohols having degrees of
20 branching between 0.9 and 2. The alcohols are prepared by homogeneously catalyzed dimerization of internal olefins and subsequent hydroformylation, where the n-proportion in the olefin to be dimerized is more than 85% by weight. A particular advantage of these alcohols is given as the cold washing behavior of their sulfates. Information about the properties of the corresponding ethoxylates and sulfates thereof is not given in this publication. A further advantage of this
25 process is given as being the fact that, for the preparation of the alcohols, no propene- or butene-containing olefin mixtures are used, but mixtures which comprise at least 85% by weight of C₆- to C₁₀-olefins.

30 In the parallel German patent application 198 59 911.0, the preparation of tridecanol mixtures by dimerization of so-called "dimer propene" and subsequent hydroformylation of the dodecene mixture is described. It typically comprises less

than 30% by weight of linear hexenes. The "dimer propene" used in this process is a hexene isomer mixture which is produced in refinery processes during the oligomerization of propene, e.g. by the DIMERSOL process (cf. Cornils/Herrmann, Applied Homogeneous Catalysis, Verlag Chemie (1996)).

5

Other suitable starting materials for this process are olefin mixtures which are prepared by dimerization of C3/C4 cuts or C4 fractions, e.g. by the DIMERSOL process. The relation between ecotoxicity and biodegradability of the alkanol mixtures prepared is, however, still not optimal.

10

From the discussed disadvantages of the prior art, in particular also from the need to reduce the preparation costs, arises the object of providing a process for the preparation of surfactant alcohols which are advantageous in terms of application, in which the use of expensive raw materials, in particular costly ethylene, can be avoided.

15

Surprisingly, we have now found that branched olefins and alcohols (oxo alcohols), which can be further processed to give very highly effective surfactants - referred to below as "surfactant alcohols" - and which have an advantageous combination of favorable properties as regards their preparation and further processing and in particular with regard to their ecotoxicity and biodegradability, can be obtained if the process is carried out according to the invention as described below.

20

This invention relates to a process for the preparation of surfactant alcohols which have particularly advantageous properties with regard to ecotoxicity and biodegradability and of corresponding surfactant alcohol ethers by

25

- a) dimerization of olefin mixtures,
- b) derivatization to give primary alcohols, and
- c) optional subsequent alkoxylation,

30

which comprises using an olefin mixture which comprises from 30 to 80% by weight of linear hexene isomers and overall at least 60% by weight of hexene isomers.

The olefin mixture used for the process according to the invention must therefore satisfy two prerequisites:

35

1. the sum of all hexene isomers present therein, i.e. linear and branched hexene isomers, must be at least 60% by weight, and

5 2. the sum of the linear hexene isomers present therein must be from 30 to 80% by weight.

10 It has been found that products with very favorable overall properties can be obtained by the process according to the invention in a particularly economical manner when an olefin mixture is used which is obtained by mixing dimer propene and linear hexenes in the weight ratio from 0.3:1 to 1:0.1.

15 For use in the process according to the invention, particular preference is given to an olefin mixture consisting of a mixture of dimer propene and linear hexenes in the weight ratio from 0.5:1 to 1:0.5.

20 In the dimerization of hexene isomer mixtures (step a) of the process according to the invention, dimerization products are obtained which, with regard to further processing to surfactant alcohols, have particularly favorable components and a particularly advantageous composition when a dimerization catalyst is used which comprises at least one element of subgroup VIII of the Periodic Table and the catalyst composition and the reaction conditions are chosen such that a dimer mixture is obtained which comprises less than 10% by weight of compounds which have a structural element of the formula I (vinylidene group)



in which A¹ and A² are aliphatic hydrocarbon radicals.

30 The dimerization can be carried out with homogeneous or heterogeneous catalysis. Preference is given to the heterogeneous procedure since with this, on the one hand, catalyst removal is simplified, making the process more economical, and on the other hand no waste waters injurious to the environment are produced, as usually form during the removal of dissolved catalysts, for example by hydrolysis.

Another advantage of the heterogeneous process is that the dimerization product does not contain halogens, in particular chlorine or fluorine. Homogeneously soluble catalysts generally contain halide-containing ligands or are used in combination with halogen-containing cocatalysts. From such catalyst systems, halogen can be incorporated into the dimerization products, which considerably adversely affects both product quality and further processing, in particular hydroformylation to give surfactant alcohols.

For the heterogeneous catalysis, use is advantageously made of combinations of oxides of metals of subgroup VIII with aluminum oxide on support materials made from silicon and titanium oxides, as are known, for example, from DE-A-43 39 713. The heterogeneous catalyst can be used in a fixed bed – then preferably in coarsely particulate form as 1 to 1.5 mm chips – or in suspended form (particle size 0.05 to 0.5 mm). In the case of a heterogeneous procedure, the dimerization is advantageously carried out at temperatures of from 80 to 200°C, preferably from 100 to 180°C, at the pressure prevailing at the reaction temperature, optionally also under a protective gas at a pressure above atmospheric, in a closed system. To achieve optimal conversions, the reaction mixture is advantageously circulated repeatedly, a certain proportion of the circulating product being continuously bled out of the system and replaced by starting material.

In the dimerization according to the invention, mixtures of monounsaturated hydrocarbons are obtained whose components predominantly have a chain length twice that of the starting olefins.

The dimerization catalysts and the reaction conditions are expediently chosen within the framework of the above details such that the olefin mixtures prepared according to the invention comprise a high proportion - as a rule greater than 85%, in particular greater than 90% - of components with branches, and a low proportion - as a rule below 15, in particular below 10% - of unbranched olefins. A further characteristic is that predominantly groups having (y-4) and (y-5) carbon atoms are bonded to the branching sites of the main chain, where y is the number of carbon atoms in the dimerized monomer. The value (y-5)=0 means that no side chains are present.

Accordingly, in C₁₂-olefin mixtures prepared according to the invention, the main chain preferably carries methyl or ethyl groups at the branching points.

5 It has also been found that the dimerization mixtures can be further derivatized particularly well when the position of the double bond satisfies certain requirements. In these advantageous olefin mixtures, the position of the double bonds relative to the branches is characterized in that the ratio of "aliphatic" hydrogen atoms to "olefinic" hydrogen atoms is in the range

$$H_{\text{aliph.}} : H_{\text{olefin.}} = 47 : 1 \text{ to } 11 : 1.$$

10

"Aliphatic" hydrogen atoms are those which are bonded to carbon atoms which do not participate in a C=C double bond (π bonding), and "olefinic" hydrogen atoms are those bonded to a carbon atom which participates in a π bond.

15 Particular preference is given to dimerization mixtures in which the ratio

$$H_{\text{aliph.}} : H_{\text{olefin.}} = 23 : 1 \text{ to } 14 : 1.$$

20 The novel olefin mixtures obtainable by the first step of the process according to the invention and having the structural features given above are likewise provided by the present invention. They are useful intermediates, in particular for the further steps of the process according to the invention, namely the preparation, described below, of branched primary alcohols and surfactants, but can also be used as starting materials in other industrial processes which start from olefins, particularly when the end products are to have improved biodegradability.

25

If the olefin mixtures according to the invention are to be used for the preparation of surfactants, then they are firstly derivatized in accordance with step b) of the process according to the invention by processes known per se to give surfactant alcohols.

30 This can be achieved in a variety of ways, which either include the direct or indirect addition of water (hydration) to the double bond, or an addition of CO and hydrogen (hydroformylation) to the C=C double bond.

35 Hydration of the olefins resulting from process step a) is advantageously carried out by direct water addition with proton catalysis. An indirect route, for example via the addition of high-percentage sulfuric acid to give an alkanol sulfonate and

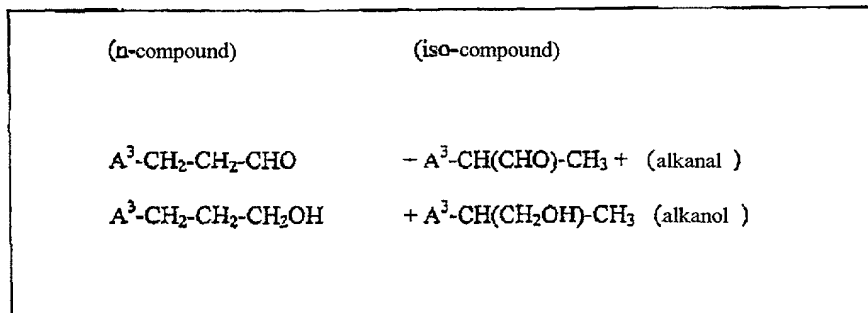
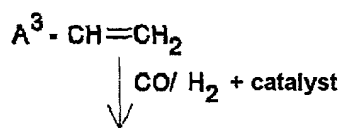
subsequent hydrolysis to give the alkanol, is, of course, also possible. The more advantageous direct water addition is carried out in the presence of acidic, in particular heterogeneous, catalysts and generally at a very high olefin partial pressure and at very low temperatures. Suitable catalysts have proven to be, in particular, phosphoric acid on supports such as, for example, SiO₂ or Celite, or else acidic ion exchangers. The choice of conditions depends on the reactivity of the olefins to be reacted and can routinely be ascertained by preliminary experiments (lit.: e.g. A.J. Kresge et al. J. Am. Chem. Soc. 93, 4907 (1971); Houben-Weyl vol. 5/4 (1960), pages 102-132 and 535-539). Hydration generally leads to mixtures of primary and secondary alkanols, in which the secondary alkanols predominate.

For the preparation of surfactants, it is more favorable to start from primary alkanols. It is therefore preferable to effect derivatization - step b) of the process according to the invention - of the olefin mixtures obtained from step a) by reaction of same with carbon monoxide and hydrogen in the presence of suitable, preferably cobalt- or rhodium-containing, catalysts to give branched primary alcohols. (Hydroformylation).

A further preferred subject-matter of the present invention is therefore a process for the preparation of mixtures of primary alkanols which are suitable, inter alia, for further processing to surfactants, by hydroformylation of olefins, which comprises using the above-described olefin mixtures according to the invention as starting material.

A good overview of the process of hydroformylation with numerous other literature references can be found, for example, in the extensive article by Beller et al. in Journal of Molecular Catalysis, A104 (1995) 17-85 or in Ullmann's Encyclopedia of Industrial Chemistry, vol. A5 (1986), page 217 et seq., page 333, and the relevant literature references.

The comprehensive information given therein allows the person skilled in the art to hydroformylate even the branched olefins according to the invention. In this reaction, CO and hydrogen are added to olefinic double bonds, giving mixtures of aldehydes and alkanols according to the following reaction equation:



(A³ = hydrocarbon radical)

- 5 The molar ratio of n- and iso-compounds in the reaction mixture is usually in the range from 1:1 to 20:1 depending on the hydroformylation process conditions chosen and the catalyst used. The hydroformylation is normally carried out in the temperature range from 90 to 200° and at a CO/H₂ pressure of from 2.5 to 35 MPa (25 to 350 bar). The mixing ratio of carbon monoxide to hydrogen depends on
- 10 whether the intention is to produce alkanals or alkanols in preference. The CO:H₂ ratio is advantageously from 10:1 to 1:10, preferably from 3:1 to 1:3, where, for the preparation of alkanals, the range of low hydrogen partial pressures is chosen, and for the preparation of alkanols the range of high hydrogen partial pressures is chosen, e.g. CO:H₂ = 1:2.
- 15 Suitable catalysts are mainly metal compounds of the formula HM(CO)₄ or M₂(CO)₈, where M is a metal atom, preferably a cobalt, rhodium or ruthenium atom.
- Generally, under hydroformylation conditions, the catalysts or catalyst precursors used in each case form catalytically active species of the formula H_xM_y(CO)_zL_q, in
- 20 which M is a metal of subgroup VIII, L is a ligand, which can be a phosphine, phosphite, amine, pyridine or any other donor compound, including in polymeric form, and q, x, y and z are integers depending on the valency and type of metal, and the covalence of the ligand L, where q can also be 0.
- The metal M is preferably cobalt, ruthenium, rhodium, palladium, platinum,
- 25 osmium or iridium and in particular cobalt, rhodium or ruthenium.

Suitable rhodium compounds or complexes are, for example, rhodium(II) and rhodium(III) salts, such as rhodium(III) chloride, rhodium(III) nitrate, rhodium(III) sulfate, potassium rhodium sulfate, rhodium(II) or rhodium(III) carboxylate, rhodium(II) and rhodium(III) acetate, rhodium (III) oxide, salts of rhodium(III) acid, such as, for example, trisammonium hexachlororhodate(III). Also suitable are rhodium complexes such as rhodium biscarbonylacetylacetonate, acetylacetonatobisethylenerrhodium(I). Preference is given to using rhodium biscarbonylacetylacetonate or rhodium acetate.

10 Suitable cobalt compounds are, for example, cobalt(II) chloride, cobalt(II) sulfate, cobalt(II) carbonate, cobalt(II) nitrate, their amine or hydrate complexes, cobalt carboxylates, such as cobalt acetate, cobalt ethylhexanoate, cobalt naphthenate, and the cobalt caprolactamate complex. Here, too, it is possible to use the carbonyl complexes of cobalt, such as dicobalt octacarbonyl, tetracobalt dodecacarbonyl and
15 hexacobalt hexadecacarbonyl.

Said compounds of cobalt, rhodium and ruthenium are known in principle and are described adequately in the literature, or they can be prepared by the person skilled in the art in a manner analogous to that for compounds already known.

20 The hydroformylation can be carried out with the addition of inert solvents or diluents or without such an addition. Suitable inert additives are, for example, acetone, methyl ethyl ketone, cyclohexanone, toluene, xylene, chlorobenzene, methylene chloride, hexane, petroleum ether, acetonitrile, and the high-boiling fractions from the hydroformylation of the dimerization products.

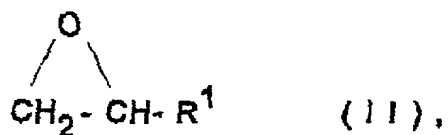
25 If the resulting hydroformylation product has too high an aldehyde content, this can be removed in a simple manner by hydrogenation, for example using hydrogen in the presence of Raney nickel or using other catalysts known for hydrogenation reactions, in particular catalysts containing copper, zinc, cobalt, nickel,
30 molybdenum, zirconium or titanium. In the process, the aldehyde fractions are largely hydrogenated to give alkanols. A virtually residue-free removal of aldehyde fractions from the reaction mixture can, if desired, be achieved by posthydrogenation, for example under particularly mild and economical conditions using an alkali metal borohydride.

35

The mixtures of branched primary alkanols, preparable by hydroformylation of the olefin mixtures according to the invention, are likewise provided by the present invention.

5 Nonionic or anionic surfactants can be prepared from the alkanols according to the invention in different ways.

Nonionic surfactants are obtained by reacting the alkanols with alkylene oxides (alkoxylation) of the formula II



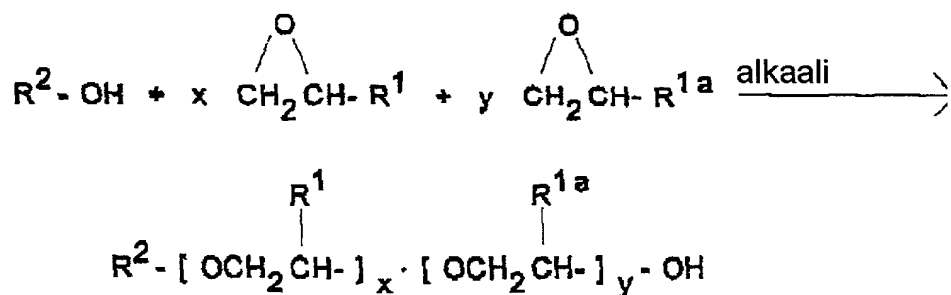
10

in which R^1 is hydrogen or a straight-chain or branched aliphatic radical of the formula $\text{C}_n\text{H}_{2n+1}$, and n is a number from 1 to 16, preferably from 1 to 8. In particular, R^1 is hydrogen, methyl or ethyl.

15 The alkanols according to the invention can be reacted with a single alkylene oxide species or with two or more different species. The reaction of the alkanols with the alkylene oxides forms compounds which in turn carry an OH group and can therefore react afresh with one molecule of alkylene oxide. Therefore, depending on the molar ratio of alkanol to alkylene oxide, reaction products are obtained which have polyether chains of varying length. The polyether chains can contain
20 from 1 to about 200 alkylene oxide structural groups. Preference is given to compounds whose polyether chains contain from 1 to 10 alkylene oxide structural groups.

The chains can consist of identical chain members, or they can have different alkylene oxide structural groups which differ from one another by virtue of their
25 radical R^1 . These various structural groups can be present within the chain in random distribution or in the form of blocks.

The reaction equation below serves to illustrate the alkoxylation of the alkanols according to the invention using, as example, a reaction with two different alkylene
30 oxides which are used in varying molar amounts x and y .



R¹ and R^{1a} are different radicals within the scope of the definitions given for R¹, and R²-OH is a branched alkanol according to the invention.

5 The alkoxylation is preferably catalyzed by strong bases, which are advantageously added in the form of an alkali metal hydroxide or alkaline earth metal hydroxide, usually in an amount of from 0.1 to 1% by weight, based on the amount of the alkanol R²-OH (cf. G. Gee et al., J. Chem. Soc. (1961), p. 1345; B. Wojtech, Makromol. Chem. 66, (1966), p. 180).

10 Acidic catalysis of the addition reaction is also possible. As well as Bronsted acids, Lewis acids, such as, for example, AlCl₃ or BF₃, are also suitable (cf. P.H. Plesch, The Chemistry of Cationic Polymerization, Pergamon Press, New York (1963)).

15 The addition reaction is carried out at temperatures of from about 120 to about 220°C, preferably from 140 to 160°C, in a sealed vessel. The alkylene oxide or the mixture of different alkylene oxides is introduced into the mixture of alkanol mixture according to the invention and alkali under the vapor pressure of the alkylene oxide mixture prevailing at the chosen reaction temperature. If desired, the alkylene oxide can be diluted by up to about 30 to 60% using an inert gas. This leads to additional security against explosive polyaddition of the alkylene oxide.

20 If an alkylene oxide mixture is used, then polyether chains are formed in which the various alkylene oxide building blocks are distributed in a virtually random manner. Variations in the distribution of the building blocks along the polyether chain arise due to varying reaction rates of the components and can also be achieved arbitrarily by continuous introduction of an alkylene oxide mixture of a
 25 program-controlled composition. If the various alkylene oxides are reacted successively, then polyether chains having block-like distribution of the alkylene oxide building blocks are obtained.

30 The length of the polyether chains varies within the reaction product in a random manner about a mean, which essentially corresponds to the stoichiometric value arising from the amount added.

099645-0100

The alkoxylates preparable starting from alkanol mixtures and olefin mixtures according to the invention are likewise provided by the present invention. They exhibit very good surface activity and can therefore be used as neutral surfactants in many areas of application.

5

Starting from the alkanol mixtures according to the invention, it is also possible to prepare surface-active glycosides and polyglycosides (oligoglycosides). These substances too have very good surfactant properties. They are obtained by single or multiple reaction (glycosidation, polyglycosidation) of the alkanol mixtures according to the invention with mono-, di- or polysaccharides with the exclusion of water and with acid catalysis. Suitable acids are, for example, HCl or H₂SO₄. As a rule, the process produces oligoglycosides having random chain length distribution, the average degree of oligomerization being from 1 to 3 saccharide radicals.

10

15

In another standard synthesis, the saccharide is firstly acetalated with acid catalysis with a low molecular weight alkanol, e.g. butanol, to give butanol glycoside. This reaction can also be carried out with aqueous solutions of the saccharide. The lower alkanol glycoside, for example butanol glycoside, is then reacted with the alkanol mixtures according to the invention to give the desired glycosides according to the invention. After the acidic catalyst has been neutralized, excess long-chain and short-chain alkanols can be removed from the equilibrium mixture, e.g. by distillation under reduced pressure.

20

25

Another standard method proceeds via the O-acetyl compounds of saccharides. The latter are converted, using hydrogen halide preferably dissolved in glacial acetic acid, into the corresponding O-acetylhalosaccharides, which react in the presence of acid-binding agents with the alkanols to give the acetylated glycosides.

30

35

Preferred for the glycosidation of the alkanol mixtures according to the invention are monosaccharides, either hexoses, such as glucose, fructose, galactose, mannose, or pentoses, such as arabinose, xylose or ribose. Particular preference for glycosidation of the alkanol mixtures according to the invention is glucose. It is, of course, also possible to use mixtures of said saccharides for the glycosidation. Glycosides having randomly distributed sugar radicals are obtained, depending on the reaction conditions. The glycosidation can also take place several times, resulting in polyglycoside chains being added to the hydroxyl groups of the alkanols. In a polyglycosidation using different saccharides, the saccharide building

blocks can be randomly distributed within the chain or form blocks of the same structural groups.

Depending on the reaction temperature chosen, furanose or pyranose structures can be obtained. To improve the solubility ratios, the reaction can also be carried out in
5 suitable solvents or diluents.

Standard processes and suitable reaction conditions have been described in various publications, for example in "Ullmann's Encyclopedia of Industrial Chemistry", 5th edition vol. A25 (1994), pages 792-793 and in the literature references given therein, by K. Igarashi, Adv. Carbohydr. Chem. Biochem. **34**, (1977), pp. 243-283,
10 by Wulff and Röhle, Angew. Chem. **86**, (1974), pp. 173-187, or in Krauch and Kunz, Reaktionen der organischen Chemie [Reactions in Organic Chemistry], pp. 405-408, Hüthig, Heidelberg, (1976).

The glycosides and polyglycosides (oligoglycosides) preparable starting from alkanol mixtures and olefin mixtures according to the invention are likewise
15 provided by the present invention.

Both the alkanol mixtures according to the invention and the polyethers prepared therefrom can be converted into anionic surfactants by esterifying (sulfating) them in a manner known per se with sulfuric acid or sulfuric acid derivatives to give
20 acidic alkyl sulfates or alkyl ether sulfates, or with phosphoric acid or its derivatives to give acidic alkyl phosphates or alkyl ether phosphates.

Sulfating reactions of alcohols have already been described, e.g. in US-A-3 462 525, 3 420 875 or 3 524 864. Details on carrying out this reaction can also be found in "Ullmann's Encyclopedia of Industrial Chemistry", 5th edition vol. A25 (1994),
25 pages 779-783 and in the literature references given therein.

If sulfuric acid itself is used for the esterification, then from 75 to 100% strength by weight, preferably from 85 to 98% strength by weight, of acid is advantageously used (so-called "concentrated sulfuric acid" or "monohydrate"). The esterification can be carried out in a solvent or diluent if one is desired for controlling the
30 reaction, e.g. the evolution of heat. In general, the alcoholic reactant is initially introduced, and the sulfating agent is gradually added with continuous mixing. If complete esterification of the alcohol component is desired, the sulfating agent and the alkanol are used in a molar ratio from 1:1 to 1:1.5, preferably from 1:1 to 1:1.2. Lesser amounts of sulfating agent can be advantageous if mixtures of alkanol
35 alkoxyates according to the invention are used and the intention is to prepare combinations of neutral and anionic surfactants. The esterification is normally

carried out at temperatures from room temperature to 85°C, preferably in the range from 45 to 75°C.

In some instances, it may be advantageous to carry out the esterification in a low-boiling water-immiscible solvent and diluent at its boiling point, the water forming during the esterification being distilled off azeotropically.

Instead of sulfuric acid of the concentration given above, for the sulfation of the alkanol mixtures according to the invention it is also possible, for example, to use sulfur trioxide, sulfur trioxide complexes, solutions of sulfur trioxide in sulfuric acid ("oleum"), chlorosulfonic acid, sulfuryl chloride and also amidosulfonic acid.

The reaction conditions are then adapted appropriately.

If sulfur trioxide is used as sulfating agent, then the reaction can also be carried out advantageously in a falling-film reactor in countercurrent, if desired also continuously.

Following esterification, the mixtures are neutralized by adding alkali and, optionally after removal of excess alkali sulfate and any solvent present, are worked up.

The acidic alkanol sulfates and alkanol ether sulfates and salts thereof obtained by sulfation of alkanols and alkanol ethers according to the invention and their mixtures are likewise provided by the present invention.

In an analogous manner, alkanols and alkanol ethers according to the invention and mixtures thereof can also be reacted (phosphated) to give acidic phosphoric esters using phosphating agents.

Suitable phosphating agents are mainly phosphoric acid, polyphosphoric acid and phosphorus pentoxide, but also POCl_3 when the remaining acid chloride functions are subsequently hydrolyzed. The phosphation of alcohols has been described, for example, in Synthesis **1985**, pages 449 to 488.

The acidic alkanol phosphates and alkanol ether phosphates obtained by phosphation of alkanols and alkanol ethers according to the invention and their mixtures are also provided by the present invention.

Finally, the use of the alkanol ether mixtures, alkanol glycosides and the acidic sulfates and phosphates of the alkanol mixtures and of the alkanol ether mixtures preparable starting from the olefin mixtures according to the invention as surfactants is also provided by the present invention.

The working examples below illustrate the preparation and use of the surfactants according to the invention.

Example 1: Heterogeneously catalyzed dimerization of hexene isomer mixtures.

A 2.5 l autoclave with rotary stirrer was charged with 200 ml of a catalyst of the following composition:

50% by weight of NiO, 34% by weight of SiO₂, 13% by weight of TiO₂, 3% by weight of Al₂O₃ (conditioned for 24 hours at 160°C in N₂, used as powder).

500 g of 1-hexene and 500 g of dimer propene, consisting of 23% by weight of n-hexenes, 70% by weight of methylpentenes and 7% by weight of dimethylbutenes, were added and reacted for 10 h at 150°C.

The resulting reaction product was worked up by distillation, and a fraction consisting of over 99% by weight of dodecenes was isolated. It has an iso index of 2.1 (4.1 methyl groups/molecule were measured following hydrogenation of the olefin by ¹H-NMR spectroscopy.), the ratio of aliphatic to olefinic protons is 14.7:1.

Example 2, hydroformylation of the dodecene mixture according to the invention.

750 g of the olefin mixture prepared as in Example 1 are hydroformylated with 2.94 g of Co₂(CO)₈ at 185°C and 280 bar of CO/H₂ (vol. ratio = 1:1) with the addition of 75 g of H₂O in a 2.5 l autoclave with lifter stirrer for 7.5 hours. Cobalt is removed from the reaction product by oxidation using 10% strength by weight acetic acid with the introduction of air at 90°C. The oxo product is hydrogenated in a 2.5 l tubular reactor in trickle mode over a Co/Mo fixed-bed catalyst at 175°C and a hydrogen pressure of 280 bar with the addition of 10% by weight of water. The alcohol mixture is worked up by distillation. The resulting tridecanol fraction has an OH number of 273 mg of KOH/g. Using ¹H-NMR spectrography, 3.5 methyl groups/molecule were detected, corresponding to a mean degree of branching of 2.5.

Example 3, fatty alcohol ethoxylate containing 7 mol of ethylene oxide.

400 g of the alkanol mixture prepared as in Example 2 are introduced with 1.5 g of NaOH into a dry 2 l autoclave. The autoclave contents are heated to 150°C, and 616 g of ethylene oxide are forced into the autoclave under pressure. After all of the ethylene oxide has been introduced into the autoclave, the autoclave is maintained at 150°C for 30 minutes. Following cooling, the catalyst is neutralized by adding sulfuric acid.

The resulting ethoxylate is a neutral surfactant. It has a cloud point of 72°C, measured in accordance with DIN 53917, 1% strength by weight in 10% strength by weight aqueous butyldiglycol solution. The surface tension of a solution of 1 g/l of the substance in water is 25.8 mN/m, measured in accordance with DIN 53914.

Example 4, fatty alcohol ethoxylate containing 3 mol of ethylene oxide.

600 g of the alkanol mixture prepared as in Example 2 are introduced with 1.5 g of NaOH into a dry 2 l autoclave. The autoclave contents are heated to 150°C, and 396 g of ethylene oxide are forced into the autoclave under pressure. After all of the ethylene oxide has been introduced into the autoclave, the autoclave is maintained at 150°C for 30 minutes. Following cooling, the catalyst is neutralized by adding sulfuric acid.

The resulting ethoxylate is a neutral surfactant. It has a cloud point of 42.1°C, measured in accordance with DIN 53917, 1% strength by weight in 10% strength by weight aqueous butyldiglycol solution. The surface tension of a solution of 1 g/l of the substance in water is 26.1 mN/m, measured in accordance with DIN 53914.

Example 5, alkyl phosphate.

300 g of the alcohol mixture prepared as in Example 2 are heated to 60°C in a stirred vessel under nitrogen, and 125 g of polyphosphoric acid are added slowly thereto. During the addition, the temperature must not exceed 65°C. Toward the end of the addition, the mixture is heated to 70°C and stirred at this temperature for a further hour.

The resulting product is an anionic surfactant. An aqueous solution of the substance in water has, at a concentration of 1 g/l, a surface tension of 28.4 mN/m, measured in accordance with DIN 53914.

Example 6, alkyl ether phosphate.

560 g of the fatty alcohol ethoxylate mixture prepared as in Example 4 are heated to 60°C in a stirred vessel under nitrogen, and 92 g of polyphosphoric acid are added slowly thereto. During the addition, the temperature must not exceed 65°C.

5 Toward the end of the addition, the mixture is heated to 70°C and stirred at this temperature for a further hour.

The resulting product is an anionic surfactant. An aqueous solution of the substance in water has, at a concentration of 1 g/l, a surface tension of 35.2 mN/m, measured in accordance with DIN 53914.

10

Example 7, alkyl sulfate.

190 g of the tridecanol mixture prepared as in Example 2 are heated to 60°C in a stirred vessel under nitrogen, and 98 g of concentrated sulfuric acid are added slowly thereto. During the addition, the temperature must not exceed 65°C. Toward

15

The resulting product is an anionic surfactant. An aqueous solution of the substance in water has, at a concentration of 1 g/l, a surface tension of 28.9 mN/m, measured in accordance with DIN 53914.

20

Example 8, alkyl ether sulfate.

480 g of the fatty alcohol ethoxylate mixture prepared as in Example 4 are heated to 60°C in a stirred vessel under nitrogen, and 146 g of concentrated sulfuric acid are added slowly thereto. During the addition, the temperature must not exceed

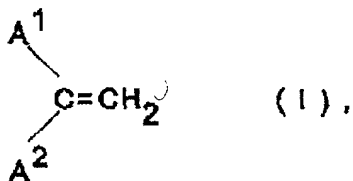
25

65°C. Toward the end of the addition, the mixture is heated to 70°C and stirred at this temperature for a further hour.

The resulting product is an anionic surfactant. An aqueous solution of the substance in water has, at a concentration of 1 g/l, a surface tension of 34.9 mN/m, measured in accordance with DIN 53914.

We claim:

1. A process for the preparation of surfactant alcohols which have particularly advantageous properties with regard to ecotoxicity and biodegradability and of corresponding surfactant alcohol ethers by
- 5 a) dimerization of olefin mixtures,
b) derivatization to give primary alcohols, and
c) optional subsequent alkoxylation,
which comprises using an olefin mixture which comprises from 30 to 80%
10 by weight of linear hexene isomers and overall at least 60% by weight of hexene isomers.
2. A process as claimed in claim 1, wherein a hexene isomer mixture is used which comprises dimer propene and linear hexenes in a weight ratio of
15 from 0.3:1 to 1:0.1.
3. A process as claimed in at least one of claims 1 and 2, wherein process step a), the dimerization, is carried out with heterogeneous catalysis.
- 20 4. A process as claimed in at least one of claims 1 to 3, wherein a dimerization catalyst is used which comprises at least one element of subgroup VIII of the Periodic Table and the catalyst composition and the reaction conditions are chosen such that a dimer mixture is obtained which comprises less than
25 10% by weight of compounds which have a structural element of the formula I (vinylidene group)



in which A¹ and A² are aliphatic hydrocarbon radicals.

5. An olefin mixture preparable by process step a) of the process of claim 1.
- 30

- 5
6. An olefin mixture as claimed in claim 5, which comprises a proportion - greater than 85%, in particular greater than 90% - of components with branches, and a low proportion - below 15, in particular below 10% - of unbranched olefins.
- 10
7. An olefin mixture as claimed in at least one of claims 5 and 6, wherein predominantly groups having (y-4) and (y-5) carbon atoms are bonded to the branching sites of the main chain, where y is the number of carbon atoms in the dimerized monomer.
- 15
8. An olefin mixture as claimed in at least one of claims 5 to 7, wherein the branched components of the dimerization mixture, in the region of 1/4 to 3/4, preferably from 1/3 to 2/3, of the chain length of the main chain, have a branch, or two branches on adjacent carbon atoms.
- 20
9. An olefin mixture as claimed in at least one of claims 5 to 8, wherein predominantly groups having one or two carbon atoms are bonded to the branching sites of the main chain.
- 25
10. An olefin mixture as claimed in at least one of claims 5 to 9, wherein, in the case of the branched components, the ratio of aliphatic to olefinic hydrogen atoms is in the range
 $H_{\text{aliph.}} : H_{\text{olefin.}} = 47 : 1 \text{ to } 11 : 1.$
- 30
11. An olefin mixture as claimed in at least one of claims 5 to 10, wherein, in the case of the branched components, the ratio of aliphatic to olefinic hydrogen atoms is in the range
 $H_{\text{aliph.}} : H_{\text{olefin.}} = 23 : 1 \text{ to } 14 : 1.$
- 35
12. A surfactant alcohol or alkoxylation products thereof, preparable by process steps a), b) and optionally c) of the process of claim 1.
13. A surfactant alcohol or alkoxylation products thereof as claimed in claim 12, which has a degree of branching between 2.0 and 3.0.

Abstract

5 The invention describes a process for the preparation of surfactant alcohols and surfactant alcohol ethers which are, inter alia, highly suitable as surfactants or for the preparation of surfactants. The process, starting from olefin mixtures having a defined minimum content of linear hexenes, gives mixtures having a predominant fraction of branched dodecenes, which are subsequently derivatized to give surfactant alcohols and then optionally alkoxyated.

10 The surfactant alcohol mixture obtained in the process has a degree of branching of from 2.0 to 3.0 and in particular exhibits advantageous properties as regards ecotoxicity and biodegradability.

A description is also given of the use of the surfactant alcohols and surfactant alcohol ethers as surfactants by glycosidation or polyglycosidation, sulfation or phosphation.

Declaration, Power of Attorney

Page 1 of 4

0050/049847

We (I), the undersigned inventor(s), hereby declare(s) that:

My residence, post office address and citizenship are as stated below next to my name,

We (I) believe that we are (I am) the original, first, and joint (sole) inventor(s) of the subject matter which is claimed and for which a patent is sought on the invention entitled

Process for the preparation of surfactant alcohols and surfactant alcohol ethers, the prepared products and their use

the specification of which

is attached hereto.

was filed on _____ as

Application Serial No. _____

and amended on _____.

was filed as PCT international application

Number _____ PCT/EP00/02416

on _____ March 17, 2000

and was amended under PCT Article 19

on _____ (if applicable).

We (I) hereby state that we (I) have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We (I) acknowledge the duty to disclose information known to be material to the patentability of this application as defined in Section 1.56 of Title 37 Code of Federal Regulations.

We (I) hereby claim foreign priority benefits under 35 U.S.C. § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed. Prior Foreign Application(s)

Application No.	Country	Day/Month/Year	Priority Claimed
19912418.3	Germany	19 March 1999	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No

We (I) hereby claim the benefit under Title 35, United States Codes, § 119(e) of any United States provisional application(s) listed below.

_____	_____
(Application Number)	(Filing Date)
_____	_____
(Application Number)	(Filing Date)

We (I) hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

Application Serial No.	Filing Date	Status (pending, patented, abandoned)
_____	_____	_____
_____	_____	_____
_____	_____	_____

And we (I) hereby appoint: Norman F. Oblon, Reg. No. 24,618; Marvin J. Spivak, Reg. No. 24,913; C. Irvin McClelland, Reg. No. 21,124; Gregory J. Maier, Reg. No. 25,599; Arthur I. Neustadt, Reg. No. 24,854; Richard D. Kelly, Reg. No. 27,757; James D. Hamilton, Reg. No. 28,421; Eckhard H. Kuesters, Reg. No. 28,870; Robert T. Pous, Reg. No. 29,099; Charles L. Gholz, Reg. No. 26,395; William E. Beaumont, Reg. No. 30,996; Jean-Paul Lavalle, Reg. No. 31,451; Stephen G. Baxter, Reg. No. 32,884; Richard L. Treanor, Reg. No. 36,379; Steven P. Weihrouch, Reg. No. 32,829; John T. Goolkasian, Reg. No. 26,142; Richard L. Chinn, Reg. No. 34,305; Steven E. Lipman, Reg. No. 30,011; Carl E. Schlier, Reg. No. 34,426; James J. Kulbaski, Reg. No. 34,648; Richard A. Neifeld, Reg. No. 35,299; J. Derek Mason, Reg. No. 35,270; Surinder Sachar, Reg. No. 34,423; Christina M. Gadiano, Reg. No. 37,628; Jeffrey B. McIntyre, Reg. No. 36,867; William T. Enos, Reg. No. 33,128; Michael E. McCabe, Jr., Reg. No. 37,182; Bradley D. Lytle, Reg. No. 40,073; and Michael R. Casey, Reg. No. 40,294; our (my) attorneys, with full powers of substitution and revocation, to prosecute this application and to transact all business in the Patent Office connected therewith; and we (I) hereby request that all correspondence regarding this application be sent to the firm of **OBLON, SPIVAK, McCLELLAND, MAIER & NEUSTADT, P. C.**, whose Post Office Address is: Fourth Floor, 1755 Jefferson Davis Highway, Arlington, Virginia 22202.

We (I) declare that all statements made herein of our (my) own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

1-00

Heiko Maas
NAME OF INVENTOR

Heiko Maas
Signature of Inventor

Date July 26, 2001

Anton-Bruckner-Str.10
67105 Schifferstadt DEX
Germany
Citizen of: Germany
Post Office Address: same as residence

2-00

Michael Röper
NAME OF INVENTOR

Michael Röper
Signature of Inventor

Date July 26, 2001

Pegauer Str.10
67157 Wachenheim DEX
Germany
Citizen of: Germany
Post Office Address: same as residence

REGISTERED

3-00

Marc Walter
NAME OF INVENTOR

Marc Walter
Signature of Inventor

Date July 26, 2001

Hans-Holbein-Str.5d
67227 Frankenthal DEX
Germany
Citizen of: France
Post Office Address: same as residence

4-00

Ralf Schulz
NAME OF INVENTOR

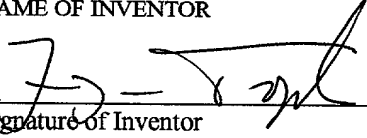
Ralf Schulz
Signature of Inventor

Date July 26, 2001

Maulbronner Hof 35
67346 Speyer DEX
Germany
Citizen of: Germany
Post Office Address: same as residence

500

Jürgen Tropsch
NAME OF INVENTOR

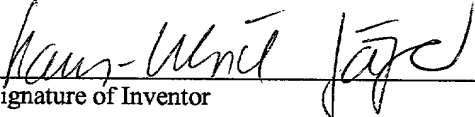

Signature of Inventor

Im oberen Berg 81
67354 Römerberg DEX
Germany
Citizen of: Germany
Post Office Address: same as residence

Date July 26, 2001

600

Hans-Ulrich Jäger
NAME OF INVENTOR


Signature of Inventor

Erschigweg 31
67434 Neustadt DEX
Germany
Citizen of: Germany
Post Office Address: same as residence

Date July 26, 2001

0050/049847